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0.22

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=> s antibiotic (3a) reporter L1 137 ANTIBIOTIC (3A) REPORTER

=> s l1 and (two hybrid or protein protein interaction)
L2 5 L1 AND (TWO HYBRID OR PROTEIN PROTEIN INTERACTION)

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PROCESSING COMPLETED FOR L2

L3 3 DUP REM L2 (2 DUPLICATES REMOVED)

=> d bib abs 1-YOU HAVE REQUESTED DATA FROM 3 ANSWERS - CONTINUE? Y/(N):y

L3 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2010 ACS on STN

AN 2003:1007706 CAPLUS

DN 140:72090

TI Expression vectors for selecting open reading frames and methods for use

IN Bradbury, Andrew R. M.; Marzari, Roberto

PA Los Alamos National Laboratory, USA

SO U.S. Pat. Appl. Publ., 25 pp. CODEN: USXXCO

DT Patent LA English

FAN.CNT 1

PATENT NO. DATE				KIND		DATE		APPLICATION NO.								
PI US 20030235814 20020619					A1		2003	1225		US 2	002-	1756	89			
					A1	A1 20031231 WO 2003-US19227										
20030618																_
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ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT AB The current invention provides method and expression vectors for selecting

open reading frames. Open reading frames present in a fragment of \mathtt{DNA}

cloned into the vectors of the invention result in creation of a fusion $% \left(1\right) =\left(1\right) +\left(1\right) +\left($

protein between the amino acid sequence encoded by the fusion protein and

a reporter protein. The vector further comprises recombination sites so

that once a recombinant that comprises an open reading frame is identified, either the reporter sequence or the open reading frame can be

removed from the vector.

L3 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2010 ACS on STN AN 1998:405969 CAPLUS

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DN
    129:77557
OREF 129:15925a,15928a
    Prokaryotic two-hybrid system for detection of
    protein-protein interactions
ΙN
    Kornacker, Michael G.
    Bristol-Myers Squibb Co., USA
PA
SO
    PCT Int. Appl., 47 pp.
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 1
    PATENT NO.
                      KIND DATE APPLICATION NO.
DATE
                       A1 19980618 WO 1997-US22703
   WO 9825947
19971210
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                E 20050630 PT 1997-952369
    PT 963376
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                T3 20050801 ES 1997-952369
    ES 2237806
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19971210

PRAI US 1996-32821P P 19961211 WO 1997-US22703 W 19971210

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT AB A two-hybrid system is provided that can detect homo-

and heterodimeric protein interactions in Escherichia coli and other

cells. This system is useful for the same applications as a yeast

two-hybrid system, i.e. interaction cloning, mapping
 protein interaction domains, analyzing protein interactions,
detecting

protein interactions, and detecting modulators thereof. The invention

concerns a prokaryotic host cell comprising: (a) a fusion protein having

(i) a first DNA-binding domain and (ii) a first interacting domain; (b) a

fusion protein having (i) a second DNA-binding domain and (ii) a second

interacting domain capable of binding to the first interacting domain; and

(c) a nucleic acid mol. having a reporter gene operatively linked to (i) a $\ \ \,$

promoter, (ii) a first operator site capable of binding to the first

DNA-binding domain, located upstream of the promoter, and (iii) a second

operator site capable of binding the second DNA-binding domain, located

downstream of the promoter of the reporter gene. Binding of the first

interacting domain to the second interacting domain is signaled by altered

expression of the reporter gene.

OSC.G 7 THERE ARE 7 CAPLUS RECORDS THAT CITE THIS RECORD (7 CITINGS)

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 3 BIOSIS COPYRIGHT (c) 2010 The Thomson Corporation on STN $\,$

DUPLICATE 1

AN 1996:434020 BIOSIS

DN PREV199699147626

TI Vectors encoding alternative antibiotic resistance for use in the yeast

two-hybrid system.

AU Watson, Michael A.; Buckholz, Richard; Weiner, Michael P. [Reprint author]

CS Dep. Mol. Sci., Glaxo Wellcome Res. Inst., Glaxo Wellcome Inc., Research

Triangle Park, NC 27709, USA

SO Biotechniques, (1996) Vol. 21, No. 2, pp. 255-259. CODEN: BTNQDO. ISSN: 0736-6205.

DT Article

LA English

ED Entered STN: 26 Sep 1996
Last Updated on STN: 26 Sep 1996

AB We have altered the antibiotic resistance of the reporter plasmids and the pJG4-5 activation-domain and pEG202 DNA binding-domain plasmids used in the Brent interaction trap/two-hybrid system. These plasmids were each previously ampicillin-resistant, resulting in an inefficient purification of any one

plasmid from a yeast strain containing all three plasmids that constitute

the complete interaction trap. By creating derivatives of each of these

plasmids expressing either kanamycin or chloramphenicol resistance, along

with the parent plasmids, we now have the option to use the interaction

trap in yeast with three ${\tt E.}$ coli differentially selectable vectors. This

will allow isolation of any one plasmid by purifying all of the
 interaction trap plasmids from yeast simultaneously and plating
E. coli

transformed with the plasmids onto the appropriate antibiotic plate to

select the particular plasmid of interest.

=> FIL STNGUIDE

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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
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-1.70		

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